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<b>(21) International Application Number:</b> PCT/EP99/09002 <b>(22) International Filing Date:</b> 23 November 1999 (23.11.99)  <b>(30) Priority Data:</b> MI98A002559      25 November 1998 (25.11.98)      IT MI99A001712      30 July 1999 (30.07.99)      IT  <b>(71) Applicant (for all designated States except US):</b> CHIESI FARMACEUTICI S.p.A. [IT/IT]; Via Palermo, 26/A, I-43100 Parma (IT).  <b>(72) Inventors; and</b> <b>(75) Inventors/Applicants (for US only):</b> LEWIS, David [GB/IT]; Chiesi Farmaceutici S.p.A., Via Palermo, 26/A, I-43100 Parma (IT). GANDERTON, David [GB/IT]; Chiesi Farmaceutici S.p.A., Via Palermo, 26/A, I-43100 Parma (IT). MEAKIN, Brian [GB/IT]; Chiesi Farmaceutici S.p.A., Via Palermo, 26/A, I-43100 Parma (IT). VENTURA, Paolo [IT/IT]; Chiesi Farmaceutici S.p.A., Via Palermo, 26/A, I-43100 Parma (IT). BRAMBILLA, Gaetano [IT/IT]; Chiesi Farmaceutici S.p.A., Via Palermo, 26/A, I-43100 Parma (IT). GARZIA, Raffaella [IT/IT]; Chiesi Farmaceutici S.p.A., Via Palermo, 26/A, I-43100 Parma (IT).		<b>(74) Agent:</b> MINOJA, Fabrizio; Bianchetti Bracco Minoja S.r.l., Via Rossini, 8, I-20122 Milan (IT).  <b>(81) Designated States:</b> AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).  <b>Published</b> <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
<b>(54) Title:</b> PRESSURISED METERED DOSE INHALERS (MDI)  <b>(57) Abstract</b> <p>The invention relates to the use of pressurised metered dose inhalers (MDIs) having part or all of their internal surfaces consisting of stainless steel, anodised aluminium or lined with an inert organic coating; and to compositions to be delivered with said MDIs.</p>		

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**"PRESSURISED METERED DOSE INHALERS (MDI) "**

The invention relates to the use of pressurised metered dose inhalers (MDIs) having part or all of their internal surfaces consisting of stainless steel, anodised aluminium or lined with an inert organic coating. The invention also relates to compositions to be delivered with said MDIs.

Pressurised metered dose inhalers are well known devices for administering pharmaceutical products to the respiratory tract by inhalation.

Active materials commonly delivered by inhalation include bronchodilators such as  $\beta_2$  agonists and anticholinergics, corticosteroids, anti-leukotrienes, anti-allergics and other materials that may be efficiently administered by inhalation, thus increasing the therapeutic index and reducing side effects of the active material.

MDI uses a propellant to expel droplets containing the pharmaceutical product to the respiratory tract as an aerosol.

For many years the preferred propellants used in aerosols for pharmaceutical use have been a group of chlorofluorocarbons which are commonly called Freons or CFCs, such as  $\text{CCl}_3\text{F}$  (Freon 11 or CFC-11),  $\text{CCl}_2\text{F}_2$  (Freon 12 or CFC-12), and  $\text{CClF}_2\text{-CClF}_2$  (Freon 114 or CFC-114).

Recently, the chlorofluorocarbon (CFC) propellants such as Freon 11 and Freon 12 have been implicated in

the destruction of the ozone layer and their production is being phased out.

Hydrofluoroalkanes [(HFAs) known also as hydrofluoro-carbons (HFCs)] contain no chlorine and are  
5 considered less destructive to ozone and these are proposed as substitutes for CFCs.

HFAs and in particular 1,1,1,2-tetrafluoroethane (HFA 134a) and 1,1,1,2,3,3,3-heptafluoropropane (HFA 227) have been acknowledged to be the best candidates  
10 for non-CFC propellants and a number of medicinal aerosol formulations using such HFA propellant systems have been disclosed.

Many of these applications, in which HFAs are used as propellant, propose the addition of one or more of  
15 adjuvants including compounds acting as co-solvents, surface active agents including fluorinated and non-fluorinated surfactants, dispersing agents including alkylpolyethoxylates and stabilizers.

In the international application n°PCT/EP98/03533  
20 filed on 10/06/98 the applicant described solution compositions for use in an aerosol inhaler, comprising an active material, a propellant containing a hydrofluoroalkane (HFA), a cosolvent and further comprising a low volatility component to increase the  
25 mass median aerodynamic diameter (MMAD) of the aerosol particles on actuation of the inhaler.

Compositions for aerosol administration via MDIs can be solutions or suspensions. Solution compositions offer several advantages: they are convenient to

manufacture being completely dissolved in the propellant vehicle and obviate physical stability problems associated with suspension compositions.

The widespread use of these formulations is limited by their chemical instability, causing the formation of degradation products.

WO94/13262 proposes the use of acids as stabilisers preventing the chemical degradation of the active ingredient in aerosol solution formulations comprising HFAs. Among the selected medicaments ipratropium bromide is comprised, for which many composition examples are supplied, in which the active ingredient is in combination with an organic or inorganic acid.

WO96/32099, WO96/32150, WO96/32151 and WO96/32345 disclose metered dose inhalers for the administration of different active ingredients in suspension in the propellant, wherein the internal surfaces of the inhaler are partially or completely coated with one or more fluorocarbon polymers optionally in combination with one or more non-fluorocarbon polymers.

Said applications do not however address the technical problem of the chemical stability of the active ingredient but they rather concern a different problem, namely that of the adhesion of micronized particles of the suspended active ingredient to the internal surfaces of the inhaler, such as the can walls, valves and sealings. It is also known from Eur. J. Pharm. Biopharm. 1997, 44, 195 that suspensions of

drugs in HFA propellant are frequently subjected to absorption of the drug particles on the valves and on the internal walls of the inhaler. The properties of an epoxy phenol resin coating of the aerosol cans have  
5 been studied to circumvent this problem.

WO 95/17195 describes aerosol compositions comprising flunisolide, ethanol and HFA propellants. It is stated in the document that conventional aerosol canisters can be used to contain the composition and  
10 that certain containers enhance its chemical and physical stability. It is suggested that the composition can be preferably contained in vials coated with resins such as epoxy resins (e.g. epoxy-phenolic resins and epoxy-urea-formaldehyde resins).

15 Actually the results reported in Tables 5, 6 and 8 respectively on pages 16 and 19 of the cited application demonstrate that flunisolide decomposes only in plastic cans (Table 8), and that the percent drug recovery in compositions stored in aluminium,  
20 glass or epoxy-phenol formaldehyde resin coated vials is practically the same (Table 8). In other words there is no difference between aluminium, glass type III or epoxy/phenol-formaldehyde resin coated aluminium vials coated by Cebal. No data are reported  
25 for other types of epoxy resins.

It has now been found that the chemical stability problems of active ingredients in solution in HFA propellants can be eliminated by storing and delivering said composition employing metered-dose

inhalers having part or all of their internal metallic surfaces consisting of stainless steel, anodised aluminium or lined with an inert organic coating.

5 The preferred material for the aerosol cans is anodised aluminium.

In the case of epoxy-phenol resin coating the choice of the suitable coating will be opportunely made on the basis of the characteristics of the active ingredient.

10 The most widely used epoxy resins in can coatings are produced by the reaction of epichlorohydrin and bisphenol A (DGEBCA). Variations in the molecular weight and in the polymerisation degree result in resins of different properties.

15 Phenoxy resins are other commercially important thermoplastic polymers derived from bisphenols and epichlorohydrin, characterized in that their molecular weights (MWs) are higher, ie, ca 45000, than those of conventional epoxy resins, ie, 8000 and lack  
20 terminal epoxide functionality.

Other multifunctional resins are epoxy-phenol-novolac and epoxy-cresol-novolac resins obtained by glycidylation of the phenol-formaldehyde (novolac) or of the o-cresol-formaldehyde (o-cresol novolac)  
25 condensates respectively.

The inhalers according to the invention effectively prevent the chemical degradation of the active ingredient.

Surprisingly and contrary to what reported in the

prior art with regard to flunisolide, we found a considerable degradation of the tested active ingredients when their formulations were stored in glass containers type III.

5        Summary of the invention

Pressurised metered dose inhalers for dispensing solution of an active ingredient in a hydrofluorocarbon propellant, a co-solvent and optionally a low-volatility component characterized in  
10 that part or all of the internal surfaces of said inhalers consist of stainless steel, anodised aluminium or are lined with an inert organic coating.

Detailed description of the invention

Pressurised metered dose inhalers are known  
15 devices, usually consisting of a main body or can, acting as a reservoir for the aerosol formulation, a cap sealing the main body and a metering valve fitted in the cap.

MDIs are usually made of a conventional material  
20 such as aluminium, tin plate, glass, plastic and the like.

According to the invention, part or all of the internal surfaces of the inhalers consists of stainless steel, anodised aluminium or is lined with  
25 an inert organic coating. One of the preferred coating consists of epoxy-phenol resin. Any kind of stainless steel may be used. Suitable epoxy-phenol resins are commercially available.

Active ingredients which may be used in the



aerosol compositions to be dispensed with the inhalers of the invention are any ingredient which can be administered by inhalation and which meets problems of chemical stability in solution in HFA propellants giving rise to a decomposition when stored in conventional materials cans and in particular in aluminium cans.

In the compositions to be delivered with the MDIs of the invention the hydrofluorocarbon propellant is preferably selected from the group of HFA 134a, HFA 227 and mixtures thereof.

The co-solvent is usually an alcohol, preferably ethanol. The low volatility component, when present, is selected from the group of glycols, particularly propylene glycol, polyethylene glycol and glycerol, alkanols such as decanol (decyl alcohol), sugar alcohols including sorbitol, mannitol, lactitol and maltitol, glycofural (tetrahydro-furfurylalcohol) and dipropylene glycol, vegetable oils, organic acids for example saturated carboxylic acids including lauric acid, myristic acid and stearic acid; unsaturated carboxylic acids including sorbic acid, and especially oleic acid; saccharine, ascorbic acid, cyclamic acid, amino acids, or aspartame, esters for example ascorbyl palmitate, isopropyl myristate and tocopherol esters; alkanes for example dodecane and octadecane; terpenes for example menthol, eucalyptol, limonene; sugars for example lactose, glucose, sucrose; polysaccharides for example ethyl cellulose, dextran; antioxidants for

example butylated hydroxytoluene, butylated hydroxyanisole; polymeric materials for example polyvinyl alcohol, polyvinyl acetate, polyvinyl pyrrolidone; amines for example ethanolamine, diethanolamine, triethanolamine; steroids for example cholesterol, cholesterol esters. The low-volatility component has a vapour pressure at 25°C lower than 0.1 kPa, preferably lower than 0.05 kPa.

The aerosols compositions to be delivered with the pressurised MDIs of the invention may contain from 0.2 to 2% by weight of said low volatility component.

Propylene glycol, polyethylene glycol, isopropyl myristate and glycerol are particularly preferred low-volatility components.

The function of the low volatility component is to modulate the MMAD of the aerosol particles. Being used at very low concentrations, it does not substantially affect the chemical stability of the compositions.

Examples of active ingredients include: anticholinergics such as ipratropium bromide, oxitropium bromide, tiotropium bromide; acetal corticosteroids such as budesonide, ciclesonide, rofleponide; chetal corticosteroids such as flunisolide, triamcinolone acetonide; other corticosteroids such as fluticasone propionate, mometasone furoate; short or long acting beta-adrenergic agonists such as salbutamol, formoterol, salmeterol, TA 2005 and their combinations. The active ingredients when possible may be present in racemic

mixtures or in form of a single enantiomer or epimer.

As said before, WO 94/13262 teaches that problems of chemical stability of medicaments and in particular of ipratropium bromide in aerosol solution compositions can be solved adding an acid, either an inorganic acid or an organic acid, to the HFA propellant/cosolvent system.

Examples of compositions containing ipratropium bromide in HFA 134a/ethanol systems further containing an inorganic acid such as hydrochloric, nitric, phosphoric or sulfuric acid or an organic acid such as ascorbic or citric acid are provided.

We found that in solution compositions comprising ipratropium bromide, a propellant containing a hydrofluoroalkane, a cosolvent and further comprising a low volatility component:

a) different decomposition rates occur with different acids: for example we found that ipratropium bromide (20 µg/dose) in a composition of 13% (w/w) ethanol, 1.0% (w/w) glycerol, 20 µl/can of 1N hydrochloric acid and HFA 134a to 12 ml/can rapidly decomposes and after 3 months storage at 40°C gives 85.0 % average of drug remaining;

b) ipratropium bromide with or without acids is stable in stainless steel, anodised aluminium or in some types of epoxy phenol resin lined cans;

c) surprisingly certain kinds of materials, such as glass, coatings proposed in the prior-art to overcome the physical absorption phenomenon of the

active ingredient, such as perfluoroalkoxyalkanes and fluorinated-ethylene-propylene polyether sulfone resins, or certain kinds of epoxy phenol coatings turned out to be completely unsatisfactory and  
5 ineffective in preventing its chemical degradation.

Another preferred active ingredient for the preparation of solution compositions in a HFA/cosolvent system to be dispensed by MDIs according to the present invention is budesonide.

10 Previously HFA/budesonide compositions have been described, in which budesonide is present in suspension in the propellant system and the composition further comprises additional ingredients such as particular kinds of surfactants (EP 504112, WO  
15 93/05765, WO 93/18746, WO 94/21229).

In WO 98/13031 it is reported that suspension formulations of budesonide have a propensity to rapidly form coarse flocs upon dispersion and redispersion which may deleteriously affect dosage  
20 reproducibility. There is also a tendency for budesonide to deposit from suspension onto the walls of the container.

To achieve stable suspensions of particulate budesonide it is employed in the prior art a  
25 composition containing a mixture of HFA propellants to match the density of the propellant mixture to be substantially identical to the density of budesonide, up to 3% of an adjuvant such as ethanol and small amounts of surfactant.

It is stated in the document that the levels of the adjuvants are low to avoid significant solubilization of drug, leading to a problem of chemical degradation and particle size increase on storage.

In the solution compositions of the present invention budesonide is chemically and physically stable.

The aerosol compositions of the invention distributed in inhalers having the internal surfaces consisting of stainless steel, anodised aluminium or coated with an inert material and preferably with epoxy-phenol resin are stable for long periods and do not undergo chemical degradation.

Also in this case a considerable degradation of the active ingredient was noticed when glass containers were used.

Analogously flunisolide and dexbudesonide (the 22R-epimer of budesonide) solutions in HFA propellant containing ethanol and a low-volatility component are stable when stored in inhalers having the internal surfaces consisting of anodised aluminium or coated with epoxy-phenol resin. Evident degradation of flunisolide was noticed when glass containers were used.

It has been also found that the low volatility component may also act as a co-solvent, thus increasing the solubility of the drug in the formulation and increasing the physical stability

and/or allowing the possibility to decrease the quantity of co-solvent required.

The following examples further illustrate the invention. In the examples and tables the different types of epoxy phenol resins are indicated with numbers in brackets corresponding to:

- (1) Epoxy-phenol lacquered aluminium vials coated by Cebal
- (2) Epoxy-phenol lacquered aluminium vials coated by Presspart
- (3) Epoxy-phenol lacquered aluminium vials coated by Nussbaum & Guhl
- (4) Epoxy-phenol lacquered aluminium vials coated by Presspart, other than (2)

Example 1

A composition containing 4.8 mg of ipratropium bromide (20  $\mu$ g/dose), 13% (w/w) ethanol, 1.0% (w/w) glycerol and HFA 134a to 12 ml/can was distributed in stainless steel, anodised aluminium, standard aluminium cans or in cans having different internal coatings and were stored at various conditions.

The results are reported in Table 1 and Table 2.

The percent drug remaining in the composition, measured by HPLC, shows that stainless steel and anodised aluminium cans as well as epoxy-phenol resins (1), (2) and (4) coated cans are effective in preventing the chemical degradation of ipratropium bromide, differently from glass cans or other tested coatings.

Example 2

The effect of different acids on the chemical stability of the composition of Example 1 was studied.

Citric, ascorbic and hydrochloric acids were added  
5 to the formulations in the amounts reported in Table 3.

The stability of the compositions was tested after 1, 2 and 5 months storage at 40°C in epoxy-phenol resin (4) coated cans.

10 Example 3

Compositions containing 12 mg of budesonide (50 µg/dose), 13% or 15% (w/w) ethanol, 1.3% (w/w) glycerol in HFA 134a to 12 ml/can were distributed in stainless steel, anodised aluminium, standard  
15 aluminium, glass cans or in cans having different internal coatings and were stored at various conditions.

The results are reported in Table 4 and 5.

The percent drug remaining in the compositions,  
20 measured by HPLC, shows the favourable effect of stainless steel, anodised aluminium and inert coating on the chemical stability of the active ingredient in respect to standard aluminium or glass cans. The best results have been obtained with stainless steel,  
25 anodised aluminium cans and with epoxy-phenol or perfluoroalkoxyalkane coatings.

Example 4

A composition containing 48 mg of dexbudesonide (200 µg/dose), 15% (w/w) ethanol, 1.3% (w/w) glycerol

in HFA 134a to 12 ml can was distributed in epoxy-phenol lacquered aluminium cans and was stored at 40°C.

5 The percent drug remaining in the composition after 8 months, measured by HPLC, was 95.4 % (average value referred to two tests).

The control of the epimeric distribution showed that there is no transfer from the 22R to the 22S epimer.

10 Example 5

Compositions containing 7.2, 12, 16.8 mg of dexbudesonide (corresponding to 30, 50 and 70 µg/dose respectively), ethanol, 0.9 (w/w) PEG 400 or isopropyl myristate (IPM) in HFA 227 to 12 ml can was distributed in aluminium anodised cans and was stored 15 70 days at 50°C. The results are reported in Table 6.

The percent drug remaining in the composition measured by HPLC shows the favourable effect of anodised aluminium cans on the chemical stability of the active ingredient. The control of the epimeric 20 distribution showed that there is no transfer from the 22R to the 22S epimer.

Example 6

The fine particle dose (FPD: weight of particles 25 having an aerodynamic diameter lower than 4.7 µm) of dexbudesonide solution compositions in HFA 134a or HFA 227, prepared following the examples 4 and 5, was determined.

The experiments were performed using the Andersen



Cascade Impactor and the data obtained are average values from 10 shots.

The results, reported in Table 7 and 8 show that dexbudesonide formulations of the invention are characterized by a very low dose and a very high fine particle dose.

The FPD gives a direct measure of the mass of particles within the specified size range and is closely related to the efficacy of the product.

#### 10      Example 7

A composition containing 60 mg of flunisolide (250 µg/dose), 15% (w/w) ethanol, 1% (w/w) glycerol in HFA 134a to 12 ml/can was distributed in anodised aluminium, glass cans or in cans having different internal coatings and were stored for 41 days at 50° C.

The results are reported in Table 9.

The percent drug remaining in the composition, measured by HPLC, shows the favourable effect of anodised aluminium and inert coating with epoxy-phenol resins on the chemical stability of the active ingredient in respect to glass cans.

#### 20      Example 8

The solubility of ipratropium bromide and micronized budesonide in ethanol, glycerol and their mixtures has been investigated.

The tests were carried out at room temperature.

a) Solubility in ethanol.

About 8.5 g of absolute ethanol were weighed into

a flask. The active ingredient (Ipratropium Bromide or Budesonide) was added in small amounts, under magnetic stirrer, until no further dissolution occurred (i.e.: a saturated solution was obtained). The flask was stirred for about 40 minutes, and left to settle overnight prior to analysis, to let the system equilibrate. The flask was kept sealed, to avoid evaporation.

The solution obtained was then filtered and tested for the amount of active ingredient, according to the conventional analytical procedure.

b) Solubility in ethanol/glycerol mixtures.

The required amounts of ethanol and glycerol were weighted into a flask, and mixed by a magnetic stirrer until a homogeneous phase was obtained.

The solubility of ipratropium bromide in ethanol is 42.48 mg/g.

The solubility data of ipratropium bromide in ethanol/glycerol mixtures are listed in Table 10.

The solubility of micronized budesonide in ethanol is 31.756 mg/g.

Solubility data of micronized budesonide in ethanol/glycerol mixtures are listed in Table 11.

The data show that both the tested active ingredients are rather soluble in ethanol, and that their solubility increases even when small percentages of glycerol are added.

The increase in solubility is maintained also in presence of HFA propellants.

TABLE 1: Percent ipratropium bromide (IPBr) recovered  
after storing the composition of Example 1  
for 8 months at 40°C in cans of different  
types

5

CAN TYPE	% RESIDUAL IPBr
Epoxy-phenol resin (4)	96
Perfluoroalkoxyalkane	57
Fluorinated-ethylene-propylene/ polyether sulphone (Xylan 8840 (R))	78
Stainless steel	96
Standard aluminium	46

15

TABLE 2: Percent ipratropium bromide (IPBr) recovered after storing the composition of Example 1 for 30 and 60 days at 50°C, or for 96 days at 40°C in cans of different types (average values referred to two tests).

CAN TYPE	% RESIDUAL IPBr			
	(% RESIDUAL IPBr RELATIVE TO			
	t=0)			
	t=0	t=30 days	t=60 days	t=96 days
		at 50°C	at 50°C	at 40°C
Epoxy phenol resin	99	89	88.5	93.5
(1)		(90)	(89.5)	(94.5)
Epoxy phenol resin	97.5	90	88.5	89
(2)		(92)	(90.5)	(91)
Epoxy phenol resin	98.5	56.5	46	52.5
(3)		(57.5)	(47)	(53.5)
Anodised aluminum	94	89	87	90.5
		(95)	(92.5)	(96.5)
Glass type III *	-	48.5	41.5	47
		(-)	(-)	(-)

\* according to Eur Pharmacopoeia 3<sup>rd</sup> Ed Suppl 1999

TABLE 3: Percent ipratropium bromide (IPBr) recovered after storing the compositions of Example 1, with different acids added, in epoxy-phenol (4) coated cans (average values referred to two tests)

	Acid	% RESIDUAL IPBr			
		(% RESIDUAL IPBr RELATIVE TO t=0)			
		t=0	t=1 month	t=2 months	t=5 months
			at 40°C	at 40°C	at 40°C
10	Citric				
	(0.6% w/w)	98	98	99	94
			(100)	(101)	(96)
	(0.3% w/w)	99	99	100	97
15			(100)	(101)	(98)
	(0.07% w/w)	99	98	99	96
			(99)	(100)	(97)
	Ascorbic	119	113	112	110
20			(95)	(94)	(92)
	Hydrochloric				
	(4 $\mu$ l-1N)	101	100	104	96
			(99)	(102)	(95)
25	(10 $\mu$ l-1N)	101	98	98	97
			(97)	(97)	(96)
	(20 $\mu$ l-1N)	100	95	98	97
			(95)	(98)	(97)
	None	97	97	98	95
			(100)	(101)	(98)

TABLE 4: Percent budesonide recovered after storing the composition of Example 3 (13% ethanol) for 7 months at 40°C in cans of different types

5

CAN TYPE	% RESIDUAL BUDESONIDE
Epoxy-phenol resin (4)	100
Fluorinated-ethylene-propylene/ polyether sulphone (Xylan 8840 (R))	93.5
Stainless steel	97
Aluminium	68
Perfluoroalkoxyalkane	100

15

TABLE 5: Percent budesonide recovered after storing the composition of Example 3 (15% ethanol) for 33 and 73 days at 50°C in cans of different types (average values referred to two tests).

CAN TYPE	% RESIDUAL BUDESONIDE (% RESIDUAL BUDESONIDE RELATIVE TO t= 0)		
	t=0	T=33 days	t=73 days
Epoxy phenol resin (1)	99.3	97.0 (97.7)	95.4 (96.1)
Epoxy phenol resin (2)	99.5	96.6 (97.0)	95.6 (96.1)
Epoxy phenol resin (3)	99.3	96.6 (97.2)	95.9 (96.5)
Anodised aluminium	99.9	99.2 (99.3)	97.7 (97.8)
Glass type III *	-	86.15 (-)	80.4 (-)

\* according to Eur Pharmacopoeia 3<sup>rd</sup> Ed Suppl 1999

These results have been confirmed storing the same formulation up to 7 months at 30°C, 40°C, 45°C and 50°C.

TABLE 6: Percent dexamethasone recovered after storing the compositions of Example 5 for 70 days at 50°C in anodised aluminium cans (average values referred to two tests).

5

Metered dose (µg)	Ethanol % (w/w)	Low vol.comp. 0.9% (w/w)	% Residual dexamethasone (% residual dexamethasone relative to t = 0)	
			t = 0 days	t = 70 days
30	5	PEG 400	95.8	95.8
				(100)
		IPM	98.1	96.8
				(98.7)
50	8	PEG 400	99.0	98.0
				(98.9)
		IPM	98.0	99.4
				(101)
70	7	PEG 400	95.7	93.75
				(98.0)
		IPM	100.4	96.3
				(96.0)

IPM = Isopropyl myristate



TABLE 7: Fine particle dose (FPD) values of  
 dexbudesonide solution formulation in HFA  
 134a containing:  
 dexbudesonide 14.4 mg/can (60 µg/shot)  
 ethanol 8 % (w/w)  
 low volatility compound 0.9% (w/w)  
 HFA 134a to 12 ml can (valve chamber volume  
 = 63 µl)  
 MMAD = 2.0 µm

Low volatility Compound	FPD (µg)	FPF (%)	Metered dose (µg)	Delivered dose (µg)
IPM	39.9	73.6	57.9	54.2
IPM	39.4	77.4	53.2	50.9

IPM = isopropyl myristate

FPF = fine particle fraction (Fine particle dose /  
 Delivered dose x 100)

FPD = weight of particles having an aerodynamic  
 diameter lower than 4.7 µm

Metered dose is given by the sum of delivered dose and  
 actuator residue.

Delivered dose is the dose delivered ex actuator.

TABLE 8: Fine particle dose (FPD) values of  
 dexbudesonide solution formulation in HFA  
 227 containing:  
 dexbudesonide 15.12 mg/can (63 µg/shot)  
 ethanol 7 % (w/w)  
 low volatility compound 0.9% (w/w)  
 HFA 227 to 12 ml can (valve chamber volume  
 = 63 µl)  
 MMAD = 2.0 µm

Low volatility Compound	FPD (µg)	FPF (%)	Metered dose (µg)	Delivered dose (µg)
IPM	45.0	75.5	63.9	59.7
PEG 400	48.5	78.9	65.5	61.5

IPM = isopropyl myristate

FPF = fine particle fraction (Fine  
 particle dose / Delivered dose x 100)

FPD = weight of particles having an  
 aerodynamic  
 diameter lower than 4.7 µm

Metered dose is given by the sum of delivered dose and  
 actuator residue

Delivered dose is the dose delivered ex actuator

TABLE 9: Percent flunisolid recovered after storing the composition of Example 7 for 41 days at 50°C in cans of different types (average values referred to two tests).

5

CAN TYPE		% RESIDUAL FLUNISOLIDE (% RESIDUAL FLUNISOLIDE RELATIVE TO t=0))		
		t=0	t=41 days	t=93 days
Epoxy	phenol	98.4	99.2	101.4
resin (1)			(101)	(103)
Epoxy	phenol	101.9	99.7	101.9
resin (2)			(97.8)	(100)
Epoxy	phenol	101.7	99.2	101.2
resin (3)			(97.5)	(99.6)
Anodised		101.6	100.4	100.7
aluminum			(98.8)	(99.1)
Glass type III *		-	-	97.5
				(-)

\* according to Eur Pharmacopoeia 3<sup>rd</sup> Ed Suppl 1999

TABLE 10: Solubility of Ipratropium Bromide in  
ethanol/glycerol mixtures

Ethanol (%)	Glycerol (%)	Ipratropium Bromide solubility (mg/g)
100	0	42.8
92.6	7.4	74.0
91.9	8.1	74.7
91.3	8.7	90.5
88.4	11.6	98.0
82.6	17.4	115.6
71.4	28.6	196.7
60	40	271.6
40	60	307.2
21.1	78.9	265.7
0	100	73.4

TABLE 11: Solubility of micronized Budesonide in ethanol/glycerol mixtures

Ethanol (%)	Glycerol (%)	Budesonide solubility (mg/g)
100	0	31.756
92.5	7.5	36.264
91.9	8.1	36.277
91.3	8.7	37.328
87.7	12.3	38.364
83.3	16.7	37.209
71.4	28.6	35.768
60	40	28.962
39.9	60.1	14.840
21.1	78.9	3.990
0	100	0.214

**CLAIMS**

1. Pressurised metered dose inhalers containing a solution of an active ingredient in a hydrofluorocarbon propellant, a co-solvent and optionally a low-volatility component characterised in that part or all of the internal surfaces of said inhalers consist of stainless steel, anodised aluminium or are lined with an inert organic coating.

2. Pressurized metered dose inhalers according to claim 1, wherein the active ingredients are selected from  $\beta_2$  agonists, steroids or anti-cholinergic agents and their combinations.

3. Pressurized metered dose inhalers according to claim 2, wherein the active ingredient is ipratropium bromide, oxitropium bromide, tiotropium bromide, flunisolide, triamcinolone acetonide, fluticasone propionate, mometasone furoate, budesonide, ciclesonide, rofleponide and epimers thereof.

4. Pressurized metered dose inhalers according to any of claims from 1 to 3, containing a low-volatility component selected from glycerol, polyethylene glycol and isopropyl myristate.

5. Pressurized metered dose inhalers according to any of claims from 1 to 4, wherein the co-solvent is ethanol.

6. Pressurized metered dose inhalers according to

any of claims from 1 to 5, wherein the propellant is selected from HFA 227, HFA 134a and their mixtures.

5 7. Pressurised metered dose inhalers according to any of claims 1 to 6 wherein the inert organic coating is perfluoroalkoxyalkane, epoxy-phenol resin or fluorinated-ethylene-propylene polyether sulfone.

10 8. Pressurised metered dose inhalers according to any of claims 1 to 7 wherein part or all of the internal surfaces are coated with an epoxy phenol resin.

9. Pressurised metered dose inhalers according to any of claims 1 to 6 wherein part or all of the internal surfaces consist of anodised aluminium.

15 10. Stabilized aerosol solution formulation consisting of an active ingredient in a hydrofluorocarbon propellant, a co-solvent and optionally a low-volatility component for use in a pressurised metered dose inhaler as claimed in any of claims 1 to 9.

20 11. Aerosol solution formulation of dexbudesonide in a hydrofluorocarbon propellant and ethanol as a co-solvent, further comprising a low volatility compound selected from glycerol, isopropylmyristate and polyethylene glycol.

# INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 99/09002

**A. CLASSIFICATION OF SUBJECT MATTER**  
IPC 7 A61K9/00

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)  
IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 96 32099 A (GLAXO WELLCOME) 17 October 1996 (1996-10-17)	1-3, 5-8, 10
Y	claims 1, 2, 4, 13, 15, 16 page 4, line 1 - line 33 page 5, line 6 - line 28 page 6, line 5 - line 8 page 6, line 21 - line 27 ---	4, 9
X	WO 95 17195 A (MINNESOTA MINING AND MANUFACTURING COMPANY) 29 June 1995 (1995-06-29) cited in the application claims 1-4, 18-20 page 18; example 29 --- -/--	1-3, 5, 6, 9

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

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"&" document member of the same patent family

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# INTERNATIONAL SEARCH REPORT

International Application No

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## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 642 992 A (CIBA-GEIGY) 15 March 1995 (1995-03-15) claim 1 column 4, line 50 - line 54 column 5, line 17 - line 18 ---	1,6
Y	WO 98 24420 A (BIOGLAN IRELAND) 11 June 1998 (1998-06-11) claims 1-4,8,9,12,17 ---	4
Y	WO 92 11236 A (SMITHKLINE BEECHAM) 9 July 1992 (1992-07-09) page 11; example 5 ---	9
A	US 4 835 145 A (PETER MAC DONALD) 30 May 1989 (1989-05-30) column 2, line 13 - line 45 column 4; examples A,B -----	11

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 99/09002

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9632099 A	17-10-1996	AU 710382 B	16-09-1999
		AU 5480996 A	30-10-1996
		BG 102021 A	31-07-1998
		BR 9604976 A	09-06-1998
		CA 2217950 A	17-10-1996
		CN 1186430 A	01-07-1998
		CZ 9703259 A	18-03-1998
		EP 0820279 A	28-01-1998
		HU 9801526 A	28-10-1998
		JP 11509433 T	24-08-1999
		NO 974737 A	11-12-1997
		NZ 306278 A	29-07-1999
		PL 322778 A	16-02-1998
		SK 138897 A	06-05-1998
WO 9517195 A	29-06-1995	AU 680967 B	14-08-1997
		AU 1098695 A	10-07-1995
		CA 2178473 A	29-06-1995
		EP 0735884 A	09-10-1996
		JP 9506896 T	08-07-1997
		NO 962585 A	18-06-1996
		NZ 276637 A	27-07-1997
		US 5980867 A	09-11-1999
		US 5776433 A	07-07-1998
EP 642992 A	15-03-1995	AT 163623 T	15-03-1998
		AU 690913 B	07-05-1998
		AU 7142994 A	09-03-1995
		CA 2130867 A	28-02-1995
		DE 59405357 D	09-04-1998
		ES 2113074 T	16-04-1998
		GR 3026507 T	31-07-1998
		JP 7076380 A	20-03-1995
WO 9824420 A	11-06-1998	AU 5402898 A	29-06-1998
		IE 80485 B	12-08-1998
		NO 992677 A	15-07-1999
		ZA 9710923 A	02-09-1998
WO 9211236 A	09-07-1992	AU 8642391 A	22-07-1992
		CA 2098298 A	20-06-1992
		EP 0563048 A	06-10-1993
		JP 6503066 T	07-04-1994
		PT 99869 A	30-11-1992
		ZA 9107468 A	30-12-1992
US 4835145 A	30-05-1989	IT 1196142 B	10-11-1988
		AT 56725 T	15-10-1990
		CA 1336513 A	01-08-1995
		DK 243085 A,B,	12-12-1985
		EP 0164636 A	18-12-1985
		ES 543499 D	01-05-1987
		ES 8705462 A	16-07-1987
		FI 852093 A,B,	12-12-1985
		JP 1588637 C	19-11-1990
		JP 2013680 B	04-04-1990
		JP 61040299 A	26-02-1986
		NO 852327 A,B,	12-12-1985

## INTERNATIONAL SEARCH REPORT

### Information on patent family members

**International Application No**

PCT/EP 99/09002

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4835145 A		US 4695625 A	22-09-1987